This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1-60. (cancelled)

61. (currently amended) A kit for assaying for the presence of a mutation associated with Familial Dysautonomia in an individual comprising primers 18F (SEQ ID NO:82) and 23R (SEQ ID NO:84) that are capable of amplifying a region of IKBKAP of sufficient size to detect a FD1 mutation at position 34,201 of SEQ ID NO:1 or a FD2 mutation at position 33,714 of SEQ ID NO:1, wherein said region amplified comprises a FD1 mutation at position 34,201 of SEQ ID NO:1 or a FD2 mutation at position 33,714 of SEQ ID NO:1.

62-67. (cancelled)

68. (previously presented) The kit of claim 61, wherein the region amplified comprises position 2,397 of SEQ ID NO:2.

69-80. (cancelled)

81. (currently amended) An isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 34,201 of SEQ ID NO:1, or the complement thereof, said oligonucleotide probe being suitable for the detection of the FD mutation at position 34,201 of SEQ ID NO:1.

82. (previously presented) The oligonucleotide probe of claim 81 which is 16 nucleotides.

83 - 86. (canceled)

- 87. (currently amended) An isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 34,201 of SEQ ID NO:1, or the complement thereof, except that the nucleotide corresponding to which is at the same position as position 34,201 of SEQ ID NO:1 is a cytosine, or a guanine in said complement, said oligonucleotide probe being suitable for the detection of the FD mutation at position 34,201 of SEQ ID NO:1.
- 88. (previously presented) The oligonucleotide probe of claim 87 which is 16 nucleotides.
- 89. (currently amended) An isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 33,714 of SEQ ID NO:1, or the complement thereof, except that the nucleotide corresponding to which is at the same position as position 33,714 of SEQ ID NO:1 is a cytosine, or a guanine in said complement, said oligonucleotide probe being suitable for the detection of the FD mutation at position 33,714 of SEQ ID NO:1.
- 90. (previously presented) The oligonucleotide probe of claim 89 which is 16 nucleotides.
- 91 99. (canceled)
- 100. (currently amended) A kit for the detection of <u>FD1 and FD2 mutation mutations</u> associated with Familial Dysautonomia in a sample from a human subject, said kit comprising <u>two different</u> isolated oligonucleotide <u>probes as follows:</u>

(1) an isolated oligonucleotide probe for the detection of the FD1 mutation selected from the group consisting of (a) through (d) below:

- (a) an isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 34,201 of SEQ ID NO:1 and being suitable for the detection of the FD mutation the FD1 mutation at position 34,201 of SEQ ID NO:1;
- (b) the complement of an isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 34,201 of SEQ ID NO:1 and being suitable for the detection of the FD mutation the FD1 mutation at position 34,201 of SEQ ID NO:1;
- (c) an isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 34,201 of SEQ ID NO:1 except that the nucleotide corresponding to which is at the same position as position 34,201 of SEQ ID NO:1 is a cytosine and being suitable for the detection of the FD mutation the FD1 mutation at position 34,201 of SEQ ID NO:1; and
- (d) the complement of an isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 34,201 of SEQ ID NO:1 except that the nucleotide corresponding to which is at the same position as position 34,201 of SEQ ID NO:1 in the complement is a guanine and being suitable for the detection of the FD mutation the FD1 mutation at position 34,201 of SEQ ID NO:1; and

Docket No. 13572.105039

- (2) an isolated oligonucleotide probe for the detection of the FD2 mutation selected from the group consisting of (e) through (h) below:
- (e) an isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 33,714 of SEQ ID NO:1 and being suitable for the detection of the FD mutation the FD2 mutation at position 33,714 of SEQ ID NO:1;
- (f) the complement of an isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 33,714 of SEQ ID NO:1 and being suitable for the detection of the FD mutation the FD2 mutation at position 33,714 of SEQ ID NO:1;
- (g) an isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 33,714 of SEQ ID NO:1 except that the nucleotide corresponding to which is at the same position as position 33,714 of SEQ ID NO:1 is a cytosine and being suitable for the detection of the FD mutation the FD2 mutation at position 33,714 of SEQ ID NO:1; and
- (h) the complement of an isolated oligonucleotide probe consisting of at least 16 contiguous nucleotides of the portion of SEQ ID NO:1 from nucleotide 32,642 to nucleotide 36,846 which include position 33,714 of SEQ ID NO:1 except that the nucleotide corresponding to which is at the same position as position 33,714 of SEQ ID NO:1 in the complement is a guanine and being suitable for the detection of the FD mutation the FD2 mutation at position 33,714 of SEQ ID NO:1.

Docket No. <u>13572.105039</u>

101. (currently amended) The <u>kit of claim 100</u>, wherein the isolated oligonucleotide probes of elaim 100 which is are each 16 nucleotides.